#### PATENT COOPERATION TREATY

From the INTERNATIONAL SEARCHING AUTHORITY To: WRITTEN OPINION OF THE see form PCT/ISA/220 INTERNATIONAL SEARCHING AUTHORITY (PCT Rule 43bis.1) Date of mailing (day/month/year) see form PCT/ISA/210 (second sheet) Applicant's or agent's file reference FOR FURTHER ACTION see form PCT/ISA/220 See paragraph 2 below International application No. International filing date (day/month/year) Priority date (day/month/year) PCT/EP2004/013427 09.12.2003 International Patent Classification (IPC) or both national classification and IPC G01N33/82 **Applicant** DSM IP ASSETS B.V. This opinion contains indications relating to the following items: Box No. I Basis of the opinion ☐ Box No. II **Priority** ☐ Box-No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability ☐ Box No. IV Lack of unity of invention Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement ☐ Box No. VI Certain documents cited Certain defects in the international application ☐ Box No. VII Box No. VIII Certain observations on the international application **FURTHER ACTION** If a demand for international preliminary examination is made, this opinion will usually be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA"). However, this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notifed the International Bureau under Rule 66.1 bis(b) that written opinions of this International Searching Authority will not be so considered. If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of three months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later. For further options, see Form PCT/ISA/220. 3. For further details, see notes to Form PCT/ISA/220. Name and mailing address of the ISA: **Authorized Officer** 

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## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/EP2004/013427

#### IAP20 Rec'd PCT/PTO 06 JUN 2006

		Box	No. I Basis of the opinion	
	1.	With the la	regard to the language, this opinion has been established on the basis of the international application in anguage in which it was filed, unless otherwise indicated under this item.	
٠.		- 1	This opinion has been established on the basis of a translation from the original language into the following language , which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)).	
	2.	. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of:		
مثر		a. typ	pe of material:	
			a sequence listing	
			table(s) related to the sequence listing	
		b. for	rmat of material:	
			in written format	
			in computer readable form	
		c. tim	ne of filing/furnishing:	
Erri Yea	٠, , , ,	÷: '□	contained in the international application as filed.	
			filed together with the international application in computer readable form.	
			furnished subsequently to this Authority for the purposes of search.	
			and the second of the contract	
<b></b> .	<b>3</b> .		In addition, in the case that more than one version or copy of a sequence listing and/or table relating theret has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.	
)	4.	Addit	ional comments:	

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International application No. PCT/EP2004/013427

Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims
1-9
No: Claims

Inventive step (IS)

Yes: Claims
1-9
No: Claims

Industrial applicability (IA)

Yes: Claims
1-9

No:

Claims

2. Citations and explanations

see separate sheet

## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (SEPARATE SHEET)

International application No. PCT/PTO 36 JUN 2006 PCT/EP2004/013427

#### Re Item V.

- 1. Reference is made to the following documents:
  - D1: HOLMBERG I ET AL: "DETERMINATION OF 25 HYDROXY VITAMIN D-3 IN SERUM BY HIGH PERFORMANCE LIQUID CHROMATOGRAPHY AND ISOTOPE DILUTION MASS SPECTROMETRY" SCANDINAVIAN JOURNAL OF CLINICAL AND LABORATORY INVESTIGATION, vol. 44, no. 4, 1984, pages 275-282, XP009052947 ISSN: 0036-5513
  - D2: LUQUE DE CASTRO M D ET AL: "Determination of vitamin D3 metabolites: State-of-the-art and trends" JOURNAL OF PHARMACEUTICAL AND BIOMEDICAL ANALYSIS, vol. 20, no. 1-2, June 1999 (1999-06), pages 1-17, XP002342738 ISSN: 0731-7085
  - D3: RYCHENER M ET AL: "A SIMPLIFIED AND IMPROVED DETERMINATION OF VITAMIN D IN FAT OIL AND MARGARINE BY HIGH-PERFORMANCE LIQUID CHROMATOGRAPHY" MITTEILUNGEN AUS DEM GEBIETE DER LEBENSMITTELUNTERSUCHUNG UND HYGIENE, vol. 76, no. 1, 1985, pages 112-124, XP009052981 ISSN: 0026-6841
  - D4: HEUDI O ET AL: "Simultaneous quantification of Vitamins A, D3 and E in fortified infant formulae by liquid chromatography-mass spectrometry" JOURNAL OF CHROMATOGRAPHY, ELSEVIER SCIENCE PUBLISHERS B.V. AMSTERDAM, NL, vol. 1022, no. 1-2, 2 January 2004 (2004-01-02), pages 115-123, XP004473612 ISSN: 0021-9673
- 2.1 Document D1 discloses (the references in parentheses applying to this document): a process for the quantitative determination of 25-hydroxy-vitamin D3 in serum comprising the steps of using directly the serum, adding tritium-labelled 25-hydroxyvitamin D3 (see scheme II), extracting with methanol/chloroform, submitting the extract to a preparative straight phase HPLC, collecting fractions and submitting them to reverse phase HPLC combined with quantification by multiple ion detection.

The differences between D1 and present claim 1 can therefore be seen in step a)

dispersing the feed sample in water and b) extracting the aqueous dispersion with tert-butyl methyl ether.

Thus, the problem solved by the present application is the provision of an alternative process for the quantification of 25-hydroxy-cholecalciferol.

D2, which is a review on the determination of vitamin D3 metabolites, discloses sample pretreatment based on (i) deproteinization steps with reagents such as acetonitrile, ammonium sulphate or ethanol or (ii) pretreatment based on saponification, which appears to be mainly used for samples containing high content of lipids such as infant formula, enriched milk, eggs, fish oil, etc followed by liquid-liquid extraction (see D2, p.6, right col. last two paragraphs and D3). D2 further comments on (iii) liquid-liquid extraction as a time-consuming method with short-comings such as loss of analyte.

D3 teaches the sample pretreatment according to D2(ii).

D4, which was published after the priority of the present document, discloses the quantification of vitamin A, D3 and E in fortified infant formulae by liquid chromatography-mass spectrometry. The food pretreatment is according to D2(ii).

Therefore, the person skilled in the art, taking into consideration the teaching of D2 and D3, would have overcome scientific prejudice by coming up with the idea of simply dispersing the feed and thereafter extracting the analyte and internal standard with tert-butyl methyl ether.

The method of claim 1 is enabled by the examples and it has been shown that the above stated problem is solved.

Therefore, the solution provided to the above stated problem as given in claim 1 involves an inventive step.

The dependent claims 2-9 add features to the process of claim 1 and thus also relate to novel and inventive subject-matter.

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International application No.

PCT/EP2004/013427